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Regult
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Listing first 45 summaries
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AAI40741	AAI16902	AAK35024	AAK09135	ABA28854	ABA60848	AAD26453	ABL03333	AAD26572	AAD26571	AAZ39405	AAZ39404	AAX89221	AAX89220	AAF44664	AAZ25100	AAF44661	AAD26573	AAF44650	AAH78263	AAS06721	ABL11987	ABL18303	ABL03417	AAF44679	AAH46902	AAH06178	AAF16067	ABA09692	ABA09608	AAK70641	AAK93262	AAK91856	AAZ99730	AAZ99731	AAV60289
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Best Local Similarity
Matches 1080; Conser
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                                                                       New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess
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23-AUG-2000;
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The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chrome and gene mapping, and in recombinant production of (II). The
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CC Note: The sequence data for this patent did not appear in the printed confictation, but was obtained in electronic format directly from WIPO CC at ftp.wipo.int/pub/published_pct_sequences.
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Best Local Similarity
Matches 1015; Conserv
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                                                       acgaagcaggaaacactggcaaatatcacatcagtgagttacgactttgatgaggaattc
                                                                                                           ccggaatttgttgctccagaaattgtgaactacgagcccctgggtctggaggctgacatg
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02-MAY-2000;
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                                                             CDNA are useful for clarifying the function of the protein encoded by the CDNA. The full length clones were obtained by construction of full length enriched cDNA libraries that were synthesised by the oligo-capping method. The primers enable the production of the full length cDNA easily without any special methods. The present sequence is a full length human cDNA of the invention.

Note: The sequence data for this patent did not form part of the printed
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                                                                                                                                                                                                                                                                                                                                                                                                                                                    P-PSDB;
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DB; AAM93338.
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su A, Sugiyama
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na T, Nagai
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K, Kojima
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S, Otsuki
                                            form part directly
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                                          (first
                      (serine/threonine kinase)
                                                                                   DNA;
    Protein
                                           entry)
                                                                                   2132
    Kinase;
                                                                                    ВP
     ZIP-kinase;
                        encoding
      serine/threonine
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Query Match
Best Local S
Matches 634
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                                                                                                         The invention provides human and murine recombinant Zipper Interacting protein Kinase (ZIP-Kinase) proteins. These proteins are serine/threonine kinases which bind the leucine zipper domain of transcription factor ATF4. Host cells containing vectors comprising the ZIP-Kinase nucleic acids are used for the recombinant expression of the proteins. ZIP-Kinase protein and DNA are useful as gene therapeutic agents against cancer, and as anti-cancer agents. The present sequence represents a DNA encoding a human ZIP kinase protein.
                                                                                                                                                                                                                                                                                                                                                      26-SEP-1997;
                                                                                                                                                                                                                                                                                                                                                                            24-SEP-1998;
                                                                                                                                                                                                                                                                                                                                                                                                  28-APR-1999
                                                                                                                                                                                                                                       New Recombinant Zipper Interacting Protein protein and DNA, useful as anticancer agent
                                                                                                                                                                                                                                                                                                         Akira S,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                   leucine zipper domain; transcription duman; murine; ss.
                                                                                                                                                                                                                 Claim
                                                                                                                                                                                                                                                                         P-PSDB;
                                                                                        Sequence 2132
                               y Match 47.6%;
Local Similarity 76.1%;
hes 634; Conservative
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atggagccattcaagcagcagaaggtggaggacttttatgacatcggagaggagctgggg
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                                     0;
                                    Score 514.6; DB 20;
Pred. No. 2.4e-132;
0; Mismatches 199;
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aagccggaaaacatcatgctgctggacaagaagtgcccaacccacgaatcaagctcatc
                                          aagcagatcctggacggcgttcactacctgcactctaagcgcatcgcacactttgacctg
                                                                                      ctctttgacttcctggcggagaaagagtcgctgacggaggacgaggccacccagttcctc
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          aagocagaaaacattatgttgttagacaagaatattcccattccacacatcaagctgatt
                                                      aagcagatootggatggggtgaactacottcacacaaagaaaattgctcactttgatotc
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RESULT AAH16158 ID AAH16158 ID AAH16158 ID AAH11 XX AAH16158 AAH11 XX AAH1615 AAH11 XX AAH1615 AAH1615
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27-AUG-1999;
11-JAN-2000;
02-MAY-2000;
09-JUN-2000;
to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combinatio of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end
                                                                                                                    The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary
                                                                                                                                                                                                                                                                                       Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   07-FEB-2001
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Sugiyama T,
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2000JP-0118776.
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2000JP-0241899.
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ı A, Nagai K
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Otsuki
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    b) a combination
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sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 1 least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs. The primers allow obtaining of the full-length cDNAs assaily without any specialised methods. AAH03166 to AAH13628 and AAH3633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893 represent human amino acid sequences; and AAH13639 to AAH13632 represent invention.
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Sequence 2224 BP; 419 A. 656 Ç 806 <u>ن</u> 343 Ŧ, 0 other;

47.68; 76.18;

Score Pred.

514.6; No. 2.4

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DB в 22; 132;

Length

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11-JAN-2000;
17-FEB-2000;
02-MAY-2000;
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Ishii S,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   New genes encoding protein kinase and protein phosphatase, useful for identifying modulators which can be used to treat human or animal disorders associated with the expression or function of these enzymes
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                                                                                                                                                                                                                                         The present sequence encodes a human protein kinase/protein phosphatase. The polypeptides are expected to participate in signal transduction in cells. The kinase phosphatases are connected with intracellular signalling pathways. Antisense oligonucleotides and compounds identified by screening (agonists or antagonists) can be used to treat human or animal disorders associated with the expression or function of the protein. In addition, the polypeptides may be used as target molecules fir drug development.
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T, Wakamatsu
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Pred. No. 2.4e
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A, Nagai K,
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2.4e-132;
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Otsuki
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T, Funaha
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                              EP911408-A2
           28-APR-1999
                                                                     murine;
                                                                                                             ZIP-kinase
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on factor AT
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kinase; cancer;

24-SEP-1998;

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        was cloned in antisense orientation into the EBV-based prival expression vector. The resulting expression library was introduced into HeLa cells. A fraction of the transfectants was selected with hygromycin B. The majority of transfected cells were selected with both hygromycin B and IFN-gamma. The cells that survived and/or grew in the presence of IFN-gamma were expanded and pooled. The extrachromosomal DNA was obtd. and cleaved with DpnI and introduced into E. coli HB101 host cells. A few bacterial clones were obtd. which included DNA antisense sequences, some of which were able to protect cells from the death-promoting effects of IFN-gamma. PCR maintified DNAs were prepd. from 10 individual bacterial clones. PCR
                                                                                                                                                                                                                                                     DNA whose expression mediates cytokine-induced programmed death - used to treat diseases or disorders associated wi uncontrolled, pathological cell growth or cytokine-induced programmed cell death.
                                                                                                                                                         DAP genes seem to play an imp. role in programmed cell death and the inhibition of their expression protects the cell from cytokine-promoted cell death. A cDNA library was generated from a mixture of mRNAs harvested after treatment of HeLa cells with IFN-gamma. It
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primers that corresp. to the immediate flanking sequence of the CDNA insertion sites in the pTK01 vector. The PCR fragments were used as labeled probes to seach Southern blots for possible cross. Thypridisation between some of the rescued antisense cDNA clones. The IO cDNA clones were classified into six distinct on or overlapping gps., some constituting several members (clones). In and some constituting a single member. Antisense cDNA clone 256 has cand some constituting a single member. Antisense cDNA clone 256 has composite sequence darked participation. The resulting composite sequence derived from 2 clones and the deduced AA sequence composite sequence derived from 2 clones and the deduced AA sequence care shown in AAQ8938 and AAR74205. The ORF is also shown in AAQ89839. Care shown in AAQ8938 and AAR74205. The ORF is also shown in AAQ89839. The care to the control of the protein composite sequence derived from 2 clones and the deduced AAQ89839. The care of the protein composite sequence derived from 2 clones and the deduced AAQ89839. The care of the protein composite sequence derived from 2 clones and the deduced AAQ89839.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 misc_feature
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            polyA_signal
                                                                                                              DNA whose expression mediates cytokine-induced programmed celdeath - used to treat diseases or disorders associated with uncontrolled, pathological cell growth or cytokine-induced programmed cell death.
                                                                                                                                                                                                                                                                                                                         12-OCT-1993;
                                                                                                                                                                                                                                                                                                                                                                                      20-APR-1995
                                                                                                                                                                                                                                                                                                                                                                                                                    WO9510630-A
               DAP genes seem to play an imp. role in programmed cell death and inhibition of their expression protects the cell from cytokine-promoted cell death. A cDNA library was generated from a mixture
                                                                                                                                                                                             P-PSDB;
                                                                                                                                                                                                                                             Kimchi A;
                                                                                                                                                                                                                                                                        (RYCU/) RYCUS
(YEDA ) YEDA 1
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                                                                               Claim 2; Fig 8; 61pp; English
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                                                                                                                                                                                               1995-178528/23.
DB; AAR74205.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                standard; cDNA;
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/label= instability
5103..5107
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/label= instability
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                was cloned in antisense orientation into the EBV-based pTK01
C expression vector. The resulting expression library was introduced into HeLB cells. A fraction of the transfectants was selected with C into HeLB cells. A fraction of the transfectants was selected with Dygromycin B. The majority of transfected cells were selected with the transfected cells were selected with Doth hygromycin B and IFN-gamma. The cells that survived and/or grew in the presence of IFN-gamma were expanded and pooled. The extrachromosomal DNA was obtd. and cleaved with DpnI and introduced into E. coli HB101 host cells. A few bacterial clones were obtd. The colinto E. coli HB101 host cells. A few bacterial clones were obtd. Which included DNA antisense sequences, some of which were able to protect cells from the death-promoting effects of IFN-gamma. C Plasmid DNAs were prepd. from 10 individual bacterial clones. PCR amplified cDNA inserts were generated from each plasmid using C primers that corresp. to the immediate flanking sequence of the cDNA insertion sites in the pTK01 vector. The PCR fragments were used as labeled probes to seach Southern blots for possible cross
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Matches
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Best Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           hybridisation between some of the rescued antisense cDNA clones. The 10 cDNA clones were classified into six distinct non-overlapping gps., some constituting several members (clones) and some constituting a single member. Antisense cDNA clone 256 has the DNA product called DAP-2. Clone 256 (DAP-2) was sequenced and used to screen a K562 lambda gtl0 cDNA library. The resulting composite sequence derived from 2 clones and the deduced AA sequence are shown in AAQ8933 and AAR74205. The ORF is also shown in AAQ89339. AAQ89338 has a poly A tail. The calculated mol. wt. of the protein is about 160 kDa. Several known domains and motifs were identified in the protein (see AAQ74205 FT).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence 5886 BP; 1447 A; 1524 C; 1500 G; 1415 T; 0
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gttgctccagaaattgtgaactacgagcccctgggtctggaggctgacatgtggagcata
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Similarity 67.5%;
59; Conservative
                                                              tggaaatgaatttaaaaacatatttgggactccagagttt
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Pred. No. 7.7e-114;
0; Mismatches 294;
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Claim 2; Fig

8; 157pp; English

e.g. cancers, auto:immune

disease or neurological disease

cell

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Alzheimer's; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Death associated protein; DAP-2; cell death; tumour cell; DAP-kinase; metastatic activity; cancer; psoriasis; autoimmune disease;
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                                                                           New isolated death associated protein nucleic acids - used for the diagnosis and treatment or disorders associated with programmed ce
                                                                                                                                                                                        WPI; 1998-520781/44.
                                                                                                                                                                                                                                              Kimchi A;
                                                                                                                                                                                                                                                                                             (YEDA ) YEDA
                                                                                                                                                                                                                                                                                                                                                  03-MAR-1997;
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The present sequence encodes a human cardiovascular system associated protein kinase-4 (CSAPK-4). CSAPK polypeptides are involved in signalling pathways associated with cell growth and differentiation. The CSAPK polypeptides and polynucleotides are used to screen for agents that specifically modulate CSAPK, which are potential therapeutic agents. They are also used for diagnosis, prognosis or monitoring of They are also used for diagnosis, prognosis or monitoring of CSAPK-related diseases, gene mapping, tissue typing and forensic diagnostication, and for treating or preventing disorders associated with aberrant CSAPK expression or activity, especially cardiovascular with aberrant CSAPK expression or activity, especially cardiovascular diseases such as congestive heart failure. They can also be used to generate pharmacogenomics. The CSAPK polynucleotide may also be used to generate
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  09-SEP-1998;
29-SEP-1998;
                                                             09-SEP-1999;
                                                                                                                                                                                                                                                                                                                                                                    Human; cardiovascular system associated protein kinase-4; CSAPK-4; signalling pathway; cell growth; cell differentiation; gene mapping; tissue typing; forenaic identification; cardiovascular disease;
                                                                                                                                                                                                                                                                                                                                                                                                                                                     cDNA encoding
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        New nucleic acid encoding cardiovascular system associated kinase, used e.g. for diagnosis, treatment and prevention cardiovascular disease
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            aagccctccacccacggaggaggagcagcacctcc 1080
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                                                   ggccggatgaggacctgaggaactgtgagagtgacactgaggaggacatcgccaggagga
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DB; AAY84323.
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Best Local Similarity
Matches 511; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                     The invention relates to primers for synthesising full length cDNA clones. 830 cDNA molecules encoding a human protein have been assoluted and nucleotide sequences of 5'- and 3'-ends of the cDNA molecules have been determined. Primers for synthesising the full length cDNA are useful for clarifying the function of the protein encoded by cDNA. The full length clones were obtained by construction of full length enriched cDNA libraries that were synthesised by the oligo-capping method. The primers enable the production of the full length cDNA easily without any special methods. The present sequence is the nucleotide sequence of the 5'-end of a cDNA provided in the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in CD-ROM format directly from EPO.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      08-JUL-1999; 99JP-0194486.
11-JAN-2000; 2000JP-0118774.
02-MAY-2000; 2000JP-0183765.
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                                                    gacatettegagaacaagaeggaegtggteeteateetggagetggtetetggeggggag
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75.6%;
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a T, Nagai K, Kojima
                                                                                                                                                                                                                                                                                                                                                                                                    219 C;
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pred. No. 1.3e-96;
0; Mismatches 162
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Otsuki T, Koga
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11-JAN-2000;
02-MAY-2000;
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The invention relates to primers for synthesising full length cDNA clones. 830 cDNA molecules encoding a human protein have been isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA molecules have been determined. Primers for synthesising the full length cDNA are useful for clarifying the function of the protein encoded by cDNA. The full length clones were obtained by construction of full length enriched cDNA libraries that were synthesised by the oligo-capping method. The primers enable the production of the full length cDNA easily without any special methods. The present sequence was used as the representative sequence from a human clone which was used in homology searches to identify the clone.
                                                                                                                                                                                                                               Ota
                                                                                                                                                                                                                                                                                                                                                                                                                                  Human cDNA clone representative sequence,
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                                                                                                                                                            Primers useful for synthesizing in genetic manipulation -
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                                                                                                                                        11; SEQ ID NO 1722;
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2000JP-0183765
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a T, Nagai
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T, Koga
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Best Local Similarity
Matches 511; Conserv
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                      Human; immune; haematopoietic; immune/haematopoietic antigen; cytostatic; gene therapy; vaccine; metastasis; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Sequence 757 BP; 176 A; 219 C; 220 G; 133 T; 9 other;
                                                                            06-NOV-2001 (first entry)
                                                                                                  AAK70641;
                                                                                                                       AAK70641 standard; DNA; 12638
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ilarity 75.6%;
Conservative
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Pred. No. 1.3e-96;
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26-JUL-2000

14-AUG-2000

12-AUG-2000

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13-AUG-2000

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15-SEP-2000

16-SEP-2000

17-SEP-2000

18-SEP-2000

18-SEP-2000
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19-MAY-2000;
07-JUN-2000;
28-JUN-2000;
14-SEP-2000;
14-SEP-2000;
21-SEP-2000;
21-SEP-2000;
25-SEP-2000;
25-SEP-2000;
26-SEP-2000;
27-SEP-2000;
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04-FEB-2000;
24-FEB-2000;
02-MAR-2000;
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17-MAR-2000;
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2000US-018974
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2000US-0199076
2000US-0198123
2000US-0205515
2000US-0214886
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2000US-0220963.
2000US-0220964.
2000US-0224518.
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2000US-0231242.

2000US-0231243.

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2000US-0231413.

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2000US-0228924.
2000US-0229287.
2000US-0229343.
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2000US-0225447.
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2000US-0225758.
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2000US-0230437.
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27-SEP-2000;
29-SEP-2000;
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20-CCT-2000;
02-CCT-2000;
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02-CCT-2000;
03-CCT-2000;
04-CCT-2000;
06-NOV-2000;
08-NOV-2000;
17-NOV-2000;
17-NO
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2000US-0241785

2000US-0241786

2000US-024187

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CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic comming acid sequences given in AAM82170 to AAM91921. (I) have cytostatic comming acid sequences given in AAM82170 to AAM91921. (I) have cytostatic comming acid sequences given in AAM82170 to AAM91921. (I) have cytostatic comming acid sequences associated with accine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and complex they may be used to treat disorders associated with decreased complex they may be used to treat disorders associated with decreased complex the patients of the activity of (I) by expressing inactive proteins or to complex the patients own production of (I). Additionally, (I) complex the patients own production of (I). Additionally, (I) proteins and polynucleotides may be used to provent the coll to express the complex and treat immune/haematopoletic-related diseases, especially connects and cancer metastases of haematopoletic-derived cells. AAK64703 connects and cancer metastases of haematopoletic-derived cells. AAK64703 connects from the present inwune/haematopoletic antigen genomic connects and cancer metastases of haematopoletic antigen genomic consequences from the present invention. AAK54942 to AAK54950 and AAM82169 consequences used in the exemplification of the present invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis - \,
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Sequence 12638 BP; 2397 A; 3383 C;
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                 3994 G; 2864 T; 0 other;
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